

<b>Sample Name</b>	AMISULPRIDE TABLETS-IP.	<b>Mfg Lic No</b>	[REDACTED]
<b>Submitted By</b>	Quality Control Advisor. (MMDSL) Shillong.	<b>Report No</b>	[REDACTED]
<b>Address</b>	DHS LaitumkhrahNHM, EastKhasi Hills Meghalaya.	<b>Receipt Date</b>	29-Apr-24
<b>Mfg By</b>	N.S.	<b>Report Date</b>	16-May-24
<b>Supplied By</b>	N.S.	<b>Ref No</b>	N.S.
<b>Batch No</b>	<b>Mfg. Date</b>	<b>Exp Date</b>	<b>Batch Size</b>
MMDSL/QC-0210	01/2024	12/2025	N.S.
			<b>Sample Qty</b>
			60 TABS.

### TEST RESULTS

Date / Period of Performance of test 29/04/2024 to 16/05/2024.

Reference to protocol :- I.P-2022.

Description :- White round biconvex film coated tablets having scored on one side.

Identification (by HPLC) :- Complies.

Average weight :- 322.7mg.

Uniformity of weight :- Within limit.

Dissolution (by UV) :- Complies (81.2% to 87.4%) (Limit:- NLT 70% +5%)

Related substance (by HPLC) :- Complies.

Assay (by HPLC) :- Each film coated tablet contains:-

Content of	Obtd./Avg.wt.	Claim	Limit		Method
			Lower	Upper	
Amisulpride	:- 198.65mg	200.0mg	180.0mg	220.0mg	IP.

NOTE:- SAMPLE CONSUMED IN TESTING.

Report: In Opinion of the undersigned, The sample as defined in the Act and the rules made there per IP.

